PATENT COOPERATION TREATY

PCT

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

(Chapter II of the Patent Cooperation Treaty)

(PCT Artcle 36 and Rule 70)

PCT04-027	FOR FURTHER A	ACTION	See Form PCT/IPEA/416		
International application No.	International filing da	te(day/month/year)	Priority date (day/month/year)		
PCT/KR2004/001243	25 MAY 2004 (2		31 MARCH 2004 (31.03.2004)		
International Patent Classification (IPC					
Applicant					
LG HOUSEHOLD & HEAL	TH CARE LTD. et	t al	016 8 12 93		
This report is the international p Authority under Article 35 and t	reliminary examination r	eport, established by thi nt according to Article 3	s International Preliminary Examining 66.		
2. This REPORT consists of a total	of 4 she	ets, including this cover	sheet.		
sheets of the de and/or sheets co Administrative I sheets which sup beyond the discles Supplemental Beb. (sent to the International containing a sequence I Box relating to Sequence I Box No. I Basis of the	nd to the International Buscription, claims and/or ontaining rectifications au instructions). Dersede earlier sheets, busoure in the international ox. al Bureau only) a total of isting and/or tables related the Listing (see Section 80 relating to the following in the section of the sectio	drawings which have be thorized by this Authority of the which this Authority of application as filed, as (indicate type and numed thereto, in electronic 22 of the Administrative	cen amended and are the basis for this report ty (see Rule 70.16 and Section 607 of the considers contain an amendment that goes indicated in item 4 of Box No. I and the considers carrier(s)), form only, as indicated in the Supplemental		
Box No. II Priority Box No. III Non-estable	ishment of opinion with	regard to novelty, inven	tive step and industrial applicability		
<u></u>	ity of invention				
Box No. V Reasoned citations an	statement under Article 3 ad explanations supporting	35(2) with regard to nov g such statement	elty, inventive step or industrial applicability;		
Box No. VI Certain do	cuments cited				
Box No. VII Certain def	Box No. VII Certain defects in the international application				
Box No. VIII Certain obs	servations on the internat	ional application			
Date of submission of the demand		Date of completion of	of this report		
31 OCTOBER 2005	(31.10.2005)		006 (20.07.2006)		
Name and mailing address of the IPEA/KR		Authorized officer			
Korean Intellectual Propert 920 Dunsan-dong, Seo-gu, Republic of Korea	y Office Daejeon 302-701.	HONG. SUNG	RAN (FILE)		
Facsimile No. 82-42-472-7140		Telephone No. 82.4	12 -181 81 16		

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

International application No.

PCT/KR2004/001243

Box N	No. I Basis of the report		
	With regard to the language, this report is based on the otherwise indicated under this item. This report is based on translations from the ori which is the language of a translation furnished international search (under Rules 12.3 an publication of the international application international preliminary examination (un	riginal language into the following d for the purposes of: nd 23.1(b)) on (under Rule 12.4)	
10 1	ith regard to the elements of the international application the receiving Office in response to an invitation under the report. the international application as originally filed/fur	ler Article 14 are referred to in this	
×	the description: pages 1-8 pages* 9 pages*	received by this Authority on received by this Authority on	as originally filed/furnished > 27/01/2006
	the claims: pages pages* pages* 10 - 12	as amended (togeth	as originally filed/furnished her with any statment) under Article 19 27/01/2006
	pages* the drawings: pages pages* pages*	received by this Authority on received by this Authority on	as originally filed/furnished
3.	the sequence listing and/or any related table(s) - so	see Supplemental Box Relating to S	sequence Listing.
	the description, pages		
4.	This report has been established as if (some of) the made, since they have been considered to go beyo (Rule 70.2(c)). the description, pages the claims, Nos. the drawings, sheets the sequence listing (specify): any table(s) related to sequence listing (specify):	ond the disclosure as filed. as indica	ated in the Supplemental Box
• If itei	m 4 applies, some or all of those sheets may be marke	æd "supersedod "	

10/594804 IAPO1 Rec'd PCT/PTO 28 SEP 2006

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

International application No.

PCT/KR2004/001243

Box No. V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

Statement			
Novelty (N)	Claims	1 - 5	YES
	Claims	None	NO
Inventive step (IS)	Claims	1 - 5	YES
	Claims	None	NO
Industrial applicability (IA)	. Claims	1 - 5	YES
•	Claims	None	NO.

2. Citations and explanations (Rule 70.7)

This examination report is done based on the amended description and claims dated on January 27th, 2006 under Article 34.

1. Reference is made to the following document:

D1: US 5705091 (The Clorox Company) 6 January 1998 cited in the application

2. Novelty and Inventive Step of Claim 1

The subject matter of claim 1 relates to a method for preparing an ester bleach activator compound [Chemical Formula 1] comprising: (A) preparing a fatty acid monoester [Chemical Formula 2]; (B) making a chloroformate [Chemical Formula 3]; and (C) reacting the chloroformate with hydroxybenzene, its derivatives, or its salts in water.

D1 is considered to represent the most relevant state of the art. It discloses the ester bleach activator of the present invention and the preparing method thereof. According to the example 1(D1), sodium 4-(2-octanoyloxy ethoxy carbonyloxy) benzenesulfonate is prepared by the following steps: (AA) preparing 2-hydroxyethyl octanoate: (BB) making chloroformate by reacting the 2-hydroxyethyl octanoate with phosgene in the presence of pyridine which is an organic base; and (C) reacting the chloroformate with anhydrous sodium 4-hydroxybenzenesulfonate.

(Continued on Supplemental Box.)

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

International application No.

PCT/KR2004/001243

Supplemental Box

In case the space in any of the preceding boxes is not sufficient. Continuation of:

Box V.

Compared with D1, the subject matter of claim 1 is directed to substitution of water for pyridine as a solvent in step(C). It cannot be considered obvious to a person skilled in the art to use water as a solvent in step (C). In addition, the effect thereof is remarkable compared with that of D1. Thus claim 1 of the present invention cannot be easily invented by a person skilled in the art.

Therefore, novelty and inventive step of the subject matter of claim 1 can be acknowledged under PCT Article 33(2) and 33(3).

3. Novelty and Inventive Step of Claims 2-5

Claims 2-5, which are dependent on claim 1, specify the fatty acid monoester of the step (A): the reaction temperature of the step (B); and the content of the water and the reaction temperature and the reaction time of the step (C). Therefore, claims 2-5 meet the requirements of PCT Article 33(2) and 33(3) with respect to novelty and inventive step.

4. Industrial Applicability

It is an objective of the present claims 1-5 to provide a method for preparing an ester bleach activator compound. There is no reason for negating the industrial applicability of this invention. Consequently, claims 1-5 appear to meet the requirements of PCT Article 33(4).

PCT/KR2004/001243 IPEA/KR 27.01.2006 10/594804 IAPO1 Rec'd PCT/PTO 28 SEP 2006

(C) Preparing decanoyloxyethoxycarbonyloxybenzenesulfonate

10.55g NaOH and 61.24g 4-hydroxybenzenesulfonic acid sodium salt dihydrate are well dissolved at 0 to 30°C in 120ml of water. Then, 73.45g of the compound (2-chlorocarbonyl oxyethyl decanoate) is added thereto, and then stirred for 2 hours at 60°C. Subsequently, it is cooled to room temperature and then filtered to remove solvent and salt, and then dried to obtain 109.43g of white solid products having the structure of the following Chemical Formula 7.

10 Chemical Formula 7

15

As shown in the above embodiment, the method of the present invention using water as solvent in the final step is simple and economical, and shows high yield compared to the prior art described in the above background art.

INDUSTRIAL APPLICABILITY

As described above, the method of the present invention is simple and economical, so it may be usefully applied to make the ester bleach activator compound expressed by the Chemical Formula 1, which has many advantages.

WHAT IS CLAIMED:

5

15

20

- 1. A method for preparing an ester bleach activator compound expressed by the following Chemical Formula 1, the method comprising:
- (A) preparing a fatty acid monoester having the structure of the following Chemical Formula 2;
- (B) making a chloroformate having the structure of the following Chemical Formula 3 by reacting the fatty acid monoester with at least one selected from the group consisting of phosgene, diphosgene and triphosgene in the presence of base; and
- 10 (C) reacting the chloroformate with hydroxybenzene, its derivatives, or its salts in water,

Chemical Formula 1

Chemical Formula 2

$$R_1 - C - O \left(CH_2CH_2 - O \right) - H$$

Chemical Formula 3

where, in the Chemical Formulas 1, 2 and 3, R₁ is a linear or branched alkyl of 1 to 19 carbon atoms, a linear or branched alkenyl of 1 to 19 carbon atoms, or a mixture of

at least two selected from them, n is an integer from 1 to 10, and L is one selected from the group having the structure of the following Chemical Formula 4,

Chemical Formula 4

5

15

$$-0 - \begin{bmatrix} R_2 Y \\ - \end{bmatrix} - \begin{bmatrix} R_2 Y \\ - \end{bmatrix} - \begin{bmatrix} R_2 Y \\ - \end{bmatrix}$$

where, in the Chemical Formula 4, R₂ is alkyl of 1 to 20 carbon atoms or alkenyl of 1 to 20 carbon atoms, Y is one selected from the group consisting of hydrogen, chlorine, bromine, SO₃M, CO₂M and OSO₂M, and M is one selected from the group consisting of hydrogen, alkaline metal ions, ammonium ion and equivalent alkali earth metal ions.

2. The method for preparing an ester bleach activator compound according to claim 1,

wherein the fatty acid monoester of the step (A) is prepared by reacting fatty acid with ethylene glycol or ethylene oxide.

- The method for preparing an ester bleach activator compound according to claim 1, wherein a reaction temperature of the step (B) is kept in the range of 10 to 40°C.
- 4. The method for preparing an ester bleach activator compound according 25 to claim 1,

wherein the content of the water is 10 to 60 wt% on the basis of the total weight

of the water, an chloroformate and hydroxybenzene, its derivatives or its salts.

5. The method for preparing an ester bleach activator compound according to claim 1, wherein a reaction temperature and a reaction time of the step (C) are respectively in the ranges of 20 to 100°C and 0.1 to 5 hours.